

Tue Dec 9 14:27:50 2003

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Page 8

958 10.4 86.7 94 14 AAQ8049
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ALIGNMENTS

RESULT 1
 ID AAQ77634
 XX AAQ77634 standard; RNA; 18 BP.

AC AAQ77634;

XX 25-MAR-2003 (updated)
 DT 02-JUN-1995 (first entry)

DE Ribonucleotide to tenascin gene consensus mRNA initiation site -9-+9.
 XX Antisense; polynucleotide; sense strand; tenascin; complementary;
 KW consensus; initiation; extracellular; glycoprotein; muscle; translation;
 KW proliferation; growth stimulatory; transcription; vascular stenosis;
 KW post-angioplasty; restenosis; cardiac hypertrophy; vascular surgery;
 KW organ transplant; ds.
 XX Synthetic.

PN W09421664-A1.
 XX 29-SEP-1994.
 PD 24-MAR-1994; 94WO-US03206.
 PF 25-MAR-1993; 93US-0037025.
 PR (TEXA-) TEXAS BIOTECHNOLOGY CORP.
 PA Denner LA, Dixon RAP, Rege AA, Stacy DL;
 PI WPI; 1994-316926/39.
 DR Synthetic anti-sense polynucleotide - hybridises to tenascin
 XX gene, useful for inhibiting vascular smooth muscle cell
 PT proliferation.
 PT Claim 5; Page 47; 64pp; English.
 PS A series of polynucleotides, either DNA (AAQ76388 and AAQ76392-400 and
 CC AAQ77614-18) or RNA (AAQ76390 and AAQ77633-46), directed against the
 CC consensus mRNA initiation site sequence (AAQ77661) for the tenascin gene.
 CC The polynucleotides are based on the degenerate sequence (AAQ76398) of
 CC the tenascin gene. Tenascin is an extracellular matrix glycoprotein
 CC consisting of six disulphide-linked subunits, each having molecular mass of
 CC 190-250 kDa. Tenascin may be important for smooth muscle cell
 CC proliferation as the protein has growth stimulatory activity. The
 CC polynucleotides can be used to inhibit transcription of the gene or
 CC translation of the mRNA encoding tenascin. The method is applicable to a
 CC number of diseases where the proliferation of smooth muscle is involved
 CC e.g. vascular stenosis, post-angioplasty restenosis and other
 CC non-angioplasty procedures such as cardiac hypertrophy, vascular surgery
 CC and organ transplant.
 CC (Updated on 25-MAR-2003 to correct PN field.)
 XX Sequence 18 BP; 2 A; 5 C; 8 G; 3 U; 0 other;

Query Match 100.0%; Score 12; DB 15; Length 18;

Best Local Similarity 83.3%; Pred. No. 2.8e+03;

Matches 10; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCATGGTGGAGG 12

Db 6 CCAUGGUGGAGG 17

RESULT 2
 AAQ77620/c
 ID AAQ77620 standard; DNA; 18 BP.

XX AAQ77620;

XX 25-MAR-2003 (updated)
 DT 01-JUN-1995 (first entry)

DE Antisense polynucleotide binds to tenascin gene consensus at -9-+9.

XX Antisense; polynucleotide; sense strand; tenascin; complementary;
 KW consensus; initiation; extracellular; glycoprotein; muscle; translation;
 KW proliferation; growth stimulatory; transcription; vascular stenosis;
 KW post-angioplasty; restenosis; cardiac hypertrophy; vascular surgery;
 KW organ transplant; ds.
 XX Synthetic.

XX Key Location/Qualifiers

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XX PD 29-SEP-1994.
XX PF 24-MAR-1994; 94WO-US03206.
XX PR 25-MAR-1993; 93US-0037025.
XX PA (TEXA-) TEXAS BIOTECHNOLOGY CORP.
XX PI Denner LA, Dixon RAF, Rege AA, Stacy DL;
XX WPI; 1994-316926/39.
XX Synthetic anti-sense polynucleotide - hybridises to tenascin
PT gene, useful for inhibiting vascular smooth muscle cell
PT proliferation.
XX Claim 10; Page 44; 64pp; English.
XX A series of antisense polynucleotides, either DNA (AAQ76388 and
CC AAQ77619-32) or RNA (AAQ76390 and AAQ77647-60) directed against the sense
CC strand of the gene encoding tenascin. The polynucleotides are based on
CC the complementary sequence (AAQ76386) of the consensus mRNA initiation
CC site sequence (AAQ77661) for the tenascin gene. Tenascin is an
CC extracellular matrix glycoprotein consisting six disulphide-linked
CC subunits, each having molecular mass of 190-250 kDa. Tenascin may be
CC important for smooth muscle cell proliferation as the protein has growth
CC stimulatory activity. The polynucleotides can be used to inhibit
CC transcription of the gene or translation of the mRNA encoding tenascin.
CC The method is applicable to a number of diseases where the proliferation
CC of smooth muscle is involved e.g. vascular stenosis, post-angioplasty
CC restenosis and other non-angioplasty procedures such as cardiac
CC hypertrophy, vascular surgery and organ transplant.
CC (Updated on 25-MAR-2003 to correct PN field.)
XX Sequence 18 BP; 3 A; 8 C; 5 G; 2 T; 0 other;
SQ Query Match 100.0%; Score 14; DB 15; Length 18;
Best Local Similarity 100.0%; Pred. No. 4.8e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCCCATGGTGGAGG 14
Db 15 CCCCATGGTGGAGG 2

RESULT 3
AAQ77648/C
XX AAQ77648 standard; RNA; 18 BP.
XX AAQ77648;
XX 25-MAR-2003 (updated)
XX 02-JUN-1995 (first entry)
XX Antisense ribonucleotide binds to tenascin gene consensus at -9-+9.
XX Antisense; polynucleotide; sense strand; tenascin; complementary;
KW consensus; initiation; extracellular; glycoprotein; muscle; translation;
KW proliferation; growth stimulatory; transcription; vascular stenosis;
KW post-angioplasty; restenosis; cardiac hypertrophy; vascular surgery;
KW organ transplant; ds.
XX Synthetic.
XX Key Location/Qualifiers
FH misc_difference 1..18
FT /*tag= a
FT /note= "phosphodiester bonds between nucleotides
FT may be replaced by phosphorothioate bonds"
XX PN
XX

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PD 29-SEP-1994.
XX 24-MAR-1994; 94WO-US03206.
XX PR 25-MAR-1993; 93US-0037025.
XX PA (TEXA-) TEXAS BIOTECHNOLOGY CORP.
XX PI Denner LA, Dixon RAF, Rege AA, Stacy DL;
XX WPI; 1994-316926/39.
XX Synthetic anti-sense polynucleotide - hybridises to tenascin
PT gene, useful for inhibiting vascular smooth muscle cell
PT proliferation.
XX Claim 10; Page 51; 64pp; English.
XX A series of antisense polynucleotides, either DNA (AAQ76388 and
CC AAQ77619-32) or RNA (AAQ76390 and AAQ77647-60) directed against the sense
CC strand of the gene encoding tenascin. The polynucleotides are based on
CC the complementary sequence (AAQ76386) of the consensus mRNA initiation
CC site sequence (AAQ77661) for the tenascin gene. Tenascin is an
CC extracellular matrix glycoprotein consisting six disulphide-linked
CC subunits, each having molecular mass of 190-250 kDa. Tenascin may be
CC important for smooth muscle cell proliferation as the protein has growth
CC stimulatory activity. The polynucleotides can be used to inhibit
CC transcription of the gene or translation of the mRNA encoding tenascin.
CC The method is applicable to a number of diseases where the proliferation
CC of smooth muscle is involved e.g. vascular stenosis, post-angioplasty
CC restenosis and other non-angioplasty procedures such as cardiac
CC hypertrophy, vascular surgery and organ transplant.
CC (Updated on 25-MAR-2003 to correct PN field.)
XX Sequence 18 BP; 3 A; 8 C; 5 G; 2 U; 0 other;
SQ Query Match 100.0%; Score 14; DB 15; Length 18;
Best Local Similarity 100.0%; Pred. No. 4.8e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCCCATGGTGGAGG 14
Db 15 CCCCATGGTGGAGG 2

RESULT 4
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XX AAQ76393 standard; DNA; 18 BP.
XX AAQ76393;
XX 25-MAR-2003 (updated)
XX 02-JUN-1995 (first entry)
XX Polynucleotide to tenascin gene consensus mRNA initiation site -9-+9.
XX Antisense; polynucleotide; sense strand; tenascin; complementary;
KW consensus; initiation; extracellular; glycoprotein; muscle; translation;
KW proliferation; growth stimulatory; transcription; vascular stenosis;
KW post-angioplasty; restenosis; cardiac hypertrophy; vascular surgery;
KW organ transplant; ds.
XX Synthetic.
XX Key Location/Qualifiers
FH misc_difference 1..18
FT /*tag= a
FT /note= "phosphodiester bonds between nucleotides
FT may be replaced by phosphorothioate bonds"
XX PN
XX

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ALIGNMENTS

RESULT 1
 ID AAO77634 standard; RNA; 18 BP.
 XX AC AAO77634;
 XX AC AAO77634;
 DT 25-MAR-2003 (updated)
 DT 02-JUN-1995 (first entry)

DE Ribonucleotide to tenascin gene consensus mRNA initiation site -9-+9.
 XX Antisense; polynucleotide; sense strand; tenascin; complementary;
 KW consensus; initiation; extracellular; glycoprotein; muscle; translation;
 KW proliferation; growth stimulatory; transcription; vascular stenosis;
 KW post-angioplasty; restenosis; cardiac hypertrophy; vascular surgery;
 KW organ transplant; ds.
 OS Synthetic.
 XX Key Location/Qualifiers
 XX Key misc_difference 1..18 /tag= a
 FT /note= "phosphodiester bonds between nucleotides
 FT may be replaced by phosphorothioate bonds"
 XX

Human bone marrow
 Probe #19198 used
 Human liver single
 Human genome-deriv
 Human single nucle
 Human DNA encoding
 Corn tassels-derive
 Plasmid primer-1 D
 Vector pEGFP-N1 DN
 Plasmid pEGFP-N1 f
 DNA region of plas
 Human pancreatic c
 A. aeolicus lumazi
 Icelandic scallop
 Example translatio
 Control sequence o
 Translation initia
 Translation initia
 Translation initia
 Human A-raf target
 ASO primer #10 to
 Human H-Ras DNAM
 Human Znf12-PC5
 Nucleic acid combi
 G protein coupled
 Human/murine chima
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 Antisense oligonuc
 B. anthracis micro
 Ribonucleotide red
 Capture oligonucle
 NOVA probe SEQ ID
 G protein coupled
 Primer OTG5015 to
 Sense PCR primer u
 Candida albicans G
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 GAPDH PCR primer K
 GAPDH house-keepin
 Nucleotide sequenc
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PN WO9421664-A1.
 XX 29-SEP-1994.
 XX 24-MAR-1994; 94WO-US03206.
 XX 25-MAR-1993; 93US-0037025.
 XX (TEXA-) TEXAS BIOTECHNOLOGY CORP.
 PA Denner LA, Dixon RAF, Rege AA, Stacy DL;
 DR WPI; 1994-316926/39.
 XX Synthetic anti-sense polynucleotide - hybridizes to tenascin
 PT gene, useful for inhibiting vascular smooth muscle cell
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 CC A series of polynucleotides, either DNA (AAQ76388 and AAQ76392-400 and
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 CC consensus mRNA initiation site sequence (AAQ77661) for the tenascin gene.
 CC The polynucleotides are based on the degenerate sequence (AAQ76386) of
 CC the tenascin gene. Tenascin is an extracellular matrix glycoprotein
 CC consisting of six disulphide-linked subunits, each having molecular mass of
 CC 190-250 kDa. Tenascin may be important for smooth muscle cell
 CC proliferation as the protein has growth stimulatory activity. The
 CC polynucleotides can be used to inhibit transcription of the gene or
 CC translation of the mRNA encoding tenascin. The method is applicable to a
 CC number of diseases where the proliferation of smooth muscle is involved
 CC e.g. vascular stenosis, post-angioplasty restenosis and other
 CC non-angioplasty procedures such as cardiac hypertrophy, vascular surgery
 CC and organ transplant.
 CC (Updated on 25-MAR-2003 to correct PN field.)
 XX Sequence 18 BP; 2 A; 5 C; 8 G; 3 U; 0 other;
 SQ Query Match 100.0%; Score 14; DB 15; Length 18;
 Best Local Similarity 85.7%; Pred. No. 3.9e+02;
 Matches 12; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GCGCCCCCATGTGG 14
 |||||
 DB 1 GCGCCCCCAUGGUGG 14
 RESULT 2
 ID AAO77620/c
 XX AAQ77620 standard; DNA; 18 BP.
 XX AC AAO77620;
 XX AC AAO77620;
 DT 25-MAR-2003 (updated)
 DT 01-JUN-1995 (first entry)
 XX Antisense polynucleotide binds to tenascin gene consensus at -9-+9.
 DE Antisense; polynucleotide; sense strand; tenascin; complementary;
 KW consensus; initiation; extracellular; glycoprotein; muscle; translation;
 KW proliferation; growth stimulatory; transcription; vascular stenosis;
 KW post-angioplasty; restenosis; cardiac hypertrophy; vascular surgery;
 KW organ transplant; ds.
 OS Synthetic.
 XX Key Location/Qualifiers
 XX Key misc_difference 1..18 /tag= a
 FT /note= "phosphodiester bonds between nucleotides
 FT may be replaced by phosphorothioate bonds"
 XX PN WO9421664-A1.

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 999 11.2 65.9 65 24 ABN54076
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ALIGNMENTS

RESULT 1
 AAQ77634
 ID AAQ77634 standard; RNA; 18 BP.

XX AC AAQ77634;
 XX
 DT 25-MAR-2003 (updated)
 DT 02-JUN-1995 (first entry)

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 KW consensus; initiation; extracellular; glycoprotein; muscle; translation;
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 KW post-angioplasty; restenosis; cardiac hypertrophy; vascular surgery;
 KW organ transplant; ds.

XX Synthetic.

XX Key Location/Qualifiers
 XX misc_difference 1..18
 FT /*tag= a
 FT /note= "phosphodiester bonds between nucleotides
 may be replaced by phosphorothioate bonds"

IRS-1 probe 80. S
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 PCR amplified DNA
 FokI/FspI contain
 Primer for preproH
 Human pancreatic c
 Empedobacter brevi
 PCR amplified DNA
 Target DNA oligonu
 Human gene signatu
 Human gene signatu
 Human reproductive
 Human gene signatu
 HCV NS3/NS4 sequen
 PCR primer used to
 Sense PCR primer #
 Hepatitis C virus
 Human p53 gene oli
 Tumour suppression
 Kozak consensus mo
 Human spliced tran
 Human spliced tran
 Human spliced tran
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 Chimeric antibody
 Breast cancer mark
 Human gene signatu
 Rat spliced transc
 Mouse spliced tran
 Mouse spliced tran

PN WO9421664-A1.

XX 29-SEP-1994.

XX 24-MAR-1994; 94WO-US03206.

XX 25-MAR-1993; 93US-0037025.

XX (TEXA-) TEXAS BIOTECHNOLOGY CORP.

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XX WPI; 1994-316926/39.

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 gene, useful for inhibiting vascular smooth muscle cell
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 CC The polynucleotides are based on the degenerate sequence (AAQ76386) of
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 CC consisting six disulphide-linked subunits, each having molecular mass of
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 CC number of diseases where the proliferation of smooth muscle is involved
 CC e.g. vascular stenosis, post-angioplasty restenosis and other
 CC non-angioplasty procedures such as cardiac hypertrophy, vascular surgery
 CC and organ transplant.
 CC (Updated on 25-MAR-2003 to correct PN field.)

XX SQ Sequence 18 BP; 2 A; 5 C; 8 G; 3 U; 0 other;

Query Match 100.0%; Score 17; DB 15; Length 18;
 Best Local Similarity 88.2%; Pred.No.56;

Matches 15; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

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 |||||:|||||

Db 1 GGCCCCCAUGGUGAGG 17

RESULT 2

AAQ77620/c

ID AAQ77620 standard; DNA; 18 BP.

XX AC AAQ77620;

XX 25-MAR-2003 (updated)

DT 01-JUN-1995 (first entry)

XX Antisense polynucleotide binds to tenascin gene consensus at -9-+9.

XX Antisense; polynucleotide; sense strand; tenascin; complementary;
 KW consensus; initiation; extracellular; glycoprotein; muscle; translation;
 KW proliferation; growth stimulatory; transcription; vascular stenosis;
 KW post-angioplasty; restenosis; cardiac hypertrophy; vascular surgery;
 KW organ transplant; ds.

XX Synthetic.

XX Key Location/Qualifiers

XX misc_difference 1..18

FT /*tag= a

FT /note= "phosphodiester bonds between nucleotides
 may be replaced by phosphorothioate bonds"

PN WO9421664-A1.